

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 0 875 947 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:

04.11.1998 Bulletin 1998/45

(51) Int. Cl.⁶: H01L 51/20

(21) Application number: 98100752.9

(22) Date of filing: 16.01.1998

(84) Designated Contracting States:

AT BE CH DE DK ES FI FR GB GR IE IT LI LU MC
NL PT SE

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: 30.04.1997 US 847138

(71) Applicant:

Hewlett-Packard Company
Palo Alto, California 94304 (US)

(72) Inventors:

- Antoniadis, Homer
Mountain View, CA 94304 (US)
- Inbasekaran, Michael
Midland, Michigan 48640 (US)
- Woo, Edmund P.
Midland, Michigan 48640 (US)

(74) Representative:

Liesegang, Roland, Dr.-Ing. et al
FORRESTER & BOEHMERT
Franz-Joseph-Strasse 38
80801 München (DE)

(54) Oxadiazole blue-emitting organic LED's

(57) An organic OLED[10] having a cathode[16] formed from a first conducting layer, an electroluminescent layer [14] including an oxadiazole compound, and an anode[12] constructed from a second conducting layer which is transparent to light generated by the electroluminescent layer. In one embodiment, an electron transport layer[15] is sandwiched between the anode[12] and electroluminescent layers. Other embodiments utilize a hole transport layer between the electro-

luminescent layer [14] and the anode[12] either with or without the electron transport layer[15]. In one embodiment, the anode[12] is constructed from a layer of indium tin oxide and a layer of a hole transport material that bonds to indium tin oxide and which has an energy band intermediate between that of indium tin oxide and that of the hole transport layer.

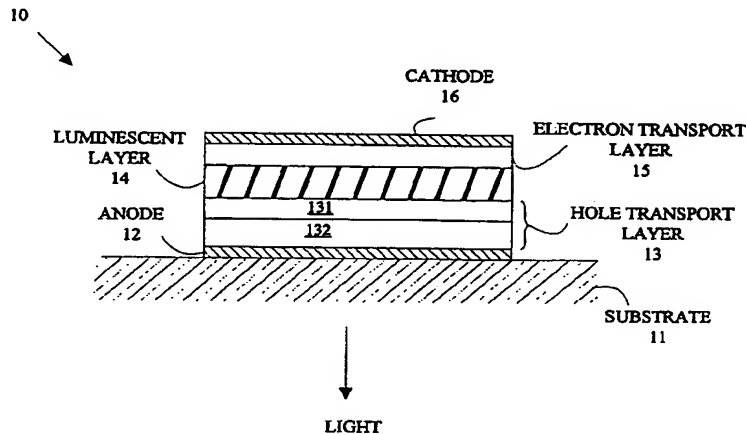


FIG. 1

EP 0 875 947 A2

Description**Field of the Invention**

5 The present invention relates to electroluminescent devices, and more particularly, to light-emitting devices based on small organic molecules.

Background of the Invention

10 Organic light-emitting diodes (OLED's) have the potential for providing inexpensive alternatives to LED's. OLED's may be fabricated by coating the appropriate surfaces with the organic material either from solution or by using conventional vacuum deposition techniques, and hence, do not require the use of high cost fabrication systems such as those utilized in the fabrication of semiconductor devices. A simple OLED may be constructed from an electroluminescent layer sandwiched between an electron injection electrode and a hole injection electrode. More complicated devices utilize electron and hole transport layers between the above mentioned electrodes and the electroluminescent layer.

15 Addressable color displays may be constructed from OLED's if individual OLED's having three primary colors can be constructed. However, compounds having a common chemical structure that can be modified with a dye to provide the primary colors and which have sufficient quantum efficiency at low operating voltages have been lacking.

Broadly, it is the object of the present invention to provide an improved OLED.

20 It is a further object of the present invention to provide a set of OLED's that can be utilized in constructing a color display.

It is a still further object of the present invention to provide an OLED that has higher quantum efficiency at low operating voltages than prior art OLED's.

25 These and other objects of the present invention will become apparent to those skilled in the art from the following detailed description of the invention and the accompanying drawings.

Summary of the Invention

30 The present invention is a organic LED having a cathode formed from a first conducting layer, an electroluminescent layer including an oxadiazole compound, and an anode constructed from a second conducting layer which is transparent to light generated by the electroluminescent layer. In one embodiment, an electron transport layer is sandwiched between the anode and electroluminescent layers. Other embodiments utilize a hole transport layer between the electroluminescent layer and the anode either with or without the electron transport layer. In one embodiment, the anode is constructed from a layer of indium tin oxide and a layer of a hole transport material that bonds to indium tin oxide and which has an energy band intermediate between that of indium tin oxide and that of the hole transport layer.

Brief Description of the Drawings

Figure 1 is a cross-sectional view of an LED according to the present invention.

40 Figures 2-3 are the chemical formulae of compounds that may be used for the electroluminescent layer of the present invention.

Figure 4 is the chemical formula of a compound that may be used for the electron transport layer of the present invention.

45 Figures 5-6 are the chemical formulae of compounds that may be used for the hole transport layer of the present invention.

Figures 7-8 are the chemical formulae of compounds that may be used between the hole transport layer and an anode constructed from indium tin oxide.

Detailed Description of the Invention

50 The present invention may be more easily understood with reference to Figure 1 which is a cross-sectional view of a OLED 10 according to the present invention. OLED 10 is typically constructed on a transparent substrate 11. A transparent anode 12 is deposited on substrate 11. Anode 12 provides a source of holes that combine with electrons injected via cathode 16 in luminescent layer 14 to generate light. To facilitate the injection of the electrons into luminescent layer 14, an optional electron transport layer 15 is utilized in some of the embodiments discussed below. Similarly, to facilitate the injection of holes into luminescent layer 14, an optional hole transport layer 13 is included between anode 12 and luminescent layer 14 in some of the embodiments discussed below. The various layers may be deposited by any of a number of conventional techniques such as spin casting, vapor deposition, etc.

The present invention is based on a luminescent layer comprising the oxadiazole compositions shown in Figures 2. Luminescent layer 14 is typically 100 to 1000 Å. In the heterocyclic compound shown in Figure 2, X is independently in each occurrence -O-, -S-, or N(R₅). Here, R₅ is a C₁₋₂₀ hydrocarbyl (which corresponds to the heterocyclic group being a 1,3,4 oxadiazole, 1,3,4-thiadiazole or 1,2,4,-triazole, respectively). R₁ is, independently in each occurrence, a C₁₋₂₀ alkyl, alkoxy, aryl, aryloxy, or a tertiary amine group -N(R₄)₂, wherein R₄ is a C₁₋₂₀ hydrocarbyl group which is optionally substituted with C₁₋₂₀ alkoxy groups. R₂ is, independently in each occurrence, H or a C₁₋₂₀ hydrocarbyl group optionally containing one or more -O-, S-, or -N(R₆)- groups, wherein R₆ is H or C₁₋₁₀ hydrocarbyl. R₃ is, independently in each occurrence, a C₁₋₂₀ hydrocarbyl group, optionally containing one or more -O-, S-, or -N(R₆)- groups; and Ar₁ is independently in each occurrence a divalent C₅₋₁₀ aromatic group.

The heterocyclic groups are preferably oxadiazole or triazole and are most preferably oxadiazoles. Each R₁ is preferably a dimethylamino or diphenylamino group. R₂ and R₃ may contain a heteroatom such as O or S, or a group of the formula: -NR₄, wherein R₄ is H or C₁₋₁₀ hydrocarbyl. Preferably, R₂ and R₃ do not contain a hydrogen atom which is active according to the Zerewitnoff test described by Kohler in *Jour. Amer. Chem. Soc.*, Vol. 49, p. 3181 (1927). Examples of suitable R₂ groups include H and C₁₋₂₀ alkyl, alkoxy, aryl, aryloxy, alkoxyaryl, and alkylcarbonyloxy groups. R₂ is preferably H. Examples of suitable R₃ groups include C₁₋₂₀ alkyl, alkoxy, aryl, aryloxy, alkoxyaryl, and alkylcarbonyloxy groups. Each R₃ is preferably selected from the group consisting of 2-ethylhexyl, n-octyl, and 4-methoxyphenyl. Examples of suitable Ar₁ groups include 1,4-phenylene, 1,3-phenylene, 2,6-naphthylene, 1,4-naphthylene, 4,4'-biphenylene, 3,5-oxazolyl, 2,5-oxadiazolyl, 2,5-triazolyl, etc.. Ar₁ is preferably a phenylene or naphthylene group, and is most preferably a 1,4- or 1,3-phenylene group.

The compounds of the present invention may be prepared by any suitable method. An example of such a method for the preparation of oxadiazole-containing compounds is by cyclization of the corresponding fluorene-containing bis-hydrazide. Methods for the cyclization of hydrazide-containing compounds are illustrated by Hayes et al. in *J. Am. Chem. Soc.*, Vol. 77, p. 1850 (1955) and by Hetzheim and Mockel, chapter on "Recent Advances in 1,3,4-Oxadiazole Chemistry" in *Advances in Heterocyclic Chemistry*, Vol. 7, pp. 183-224 (1966).

Triazole-containing compounds may be prepared by reacting imidoyl chlorides with an aromatic amine, as illustrated by E. Klingsberg in *J. Am. Chem. Soc.*, Vol. 80, pp.5786-5789 (1958). Imidoyl chlorides may be prepared by reacting bis-hydrazides with PCl₅. Thiadiazole-containing compounds may be prepared by reacting bis-hydrazides with phosphorous pentasulfide, as illustrated in *Advances in Heterocyclic Chemistry*, Vol. 9, p. 169 (1968).

Bis-hydrazides may be prepared by reacting a fluorene-containing diacid chloride with two or more equivalents of a substituted aromatic hydrazide in a dipolar, aprotic solvent such as N-methylpyrrolidone (NMP) in the presence of a base such as pyridine or triethylamine, as illustrated by F. N. Hayes et al. in *J. Am. Chem. Soc.*, Vol. 77, pp. 1850-1852 (1955). Alternatively, the reaction may also be carried out in the absence of the base wherein the solvent, NMP in particular, acts also as the base to scavenge the hydrogen halide produced in the reaction, as illustrated by A. H. Frazer and F. T. Wallenberger in *Journal of Polymer Science: Part A*, Vol. 2, pp. 1137-1145 (1964).

The fluorene-containing diacid chloride may be prepared from the corresponding diacid by any suitable technique such as, for example, reacting the diacid with thionyl chloride, oxalyl chloride, or phosgene. The fluorene-containing diacid may be prepared by reacting a dihalofluorene compound with magnesium to form a di-Grignard intermediate, which is then treated with excess CO₂ to form the desired diacid (methods for the preparation of dicarboxylic acids from haloaromatic compounds are illustrated by Gilman et al. in *Organic Synthesis, Collective*, Vol. 2, pp. 425 (1949)).

In the preferred embodiment of the present invention, luminescent layer 14 may also be constructed from luminescent layer comprising of a heterocyclic compound having the composition shown in Figure 3. In this compound, X, R₁, R₃ and Ar₁ are the groups described earlier for the compound shown in Figure 2.

In the preferred embodiment of the present invention, anode 12 is preferably constructed from transparent conducting film with a work function between 4.5 and 5.5 eV. Examples are indium tin oxide and gold. The cathode is preferably constructed from a conducting metal with a work function between 2.5 and 4.5 eV. Examples are lithium, calcium, magnesium, indium silver, aluminum or alloys of the above.

If cathode 16 is constructed from Ca, the device will generate light in the blue region of the spectrum but with low quantum efficiency. However, a less reactive cathode material is preferred. For example, cathode 16 may be constructed from an alloy of Mg/Al, Mg/Ag or Li/Al however without improvement in device efficiency. An electron transport layer such as layer 15 is preferably used to improve the efficiency of electron injection into luminescent layer 14. Electron transport layer 15 can be constructed from any compound having an energy band intermediate between that of cathode 16 and luminescent layer 14. In the preferred embodiment of the present invention, electron transport layer 15 comprises a layer of a metal chelated oxinoid compound and most preferably of tris(8-quinoline)Aluminum(Alq₃) whose formula is shown in Figure 4. The thickness of electron transport layer 15 is typically between 50 and 1000 Å.

Similarly, the hole transport layer matches luminescent layer 14 to anode 12. When anode 12 is constructed from indium tin oxide, the hole transport layer is preferably constructed from two layers. Layer 131 improves the efficiency of hole injection into luminescent layer 14 by providing a material that has an energy band between those of anode 12 and luminescent layer 14. In the preferred embodiment of the present invention, layer 131 comprises an aromatic tertiary

amine and most preferably of (N,N'-diphenyl-N,N'-bis-(3-methylphenyl)-[1,1'-biphenyl]-4,4'-diamine) (TPD) or 4,4'-Bis[N-(1-naphthyl)-N-phenylamino]-biphenyl (NPD) whose formula is shown in Figure 5 and 6 respectively.

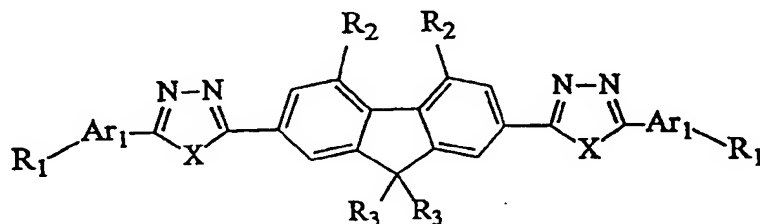
The second layer provides a physical interface between layer 131 and anode 12. It is found experimentally that compounds such as TPD do not bond well to indium tin oxide. Hence, an interface layer 132 which bonds well to indium tin oxide and has an ionization potential between that of indium tin oxide and layer 131 is utilized. In the preferred embodiment of the present invention, layer 132 is constructed from star-burst amines and most preferably from 4,4,4'-tris[N-(3-methoxyphenyl)-N-phenylamino] triphenylamine (3MeODATA) whose chemical formula is shown in Figure 7. Alternatively layer 132 is constructed from porphyrinic compounds and more preferably from copper phthalocyanine whose chemical formula is shown in Figure 8. In general, layer 131 may be viewed as being part of the anode.

Various modifications to the present invention will become apparent to those skilled in the art from the foregoing description and accompanying drawings. Accordingly, the present invention is to be limited solely by the scope of the following claims.

Claims

1. An OLED comprising: a cathode (16) comprising a first conducting layer; an electroluminescent layer (14) comprising an oxadiazole compound; and an anode (12) comprising a second conducting layer, wherein said anode (12) is transparent to light generated by said electroluminescent layer.

2. The OLED (10) of claim 1 wherein said oxadiazole compound has the formula



wherein X is independently in each occurrence -O-, -S-, or N(R₅), wherein R₅ is a C₁₋₂₀ hydrocarbonyl (which correspond to the heterocyclic group being a 1,3,4 oxadiazole, 1,3,4-thiadiazole or 1,2,4-triazole, respectively); R₁ is independently in each occurrence C₁₋₂₀ alkyl, alkoxy, aryl, aryloxy, or a tertiary amine group-N(R₄)₂, wherein R₄ is a C₁₋₂₀ hydrocarbonyl group which is optionally substituted with C₁₋₂₀ alkoxy groups; R₂ is independently in each occurrence H or a C₁₋₂₀ hydrocarbonyl group optionally containing one or more -O-, -S-, or -N(R₆)-groups, wherein R₆ is H or C₁₋₁₀ hydrocarbonyl; R₃ is independently in each occurrence a C₁₋₂₀ hydrocarbonyl group, optionally containing one or more -O-, -S-, or -N(R₆)-groups; and Ar₁ is independently in each occurrence a divalent C₅₋₁₀ aromatic group.

3. The OLED (10) of claim 2 wherein X is -O-, -S- or -N(R₅)- in each occurrence.

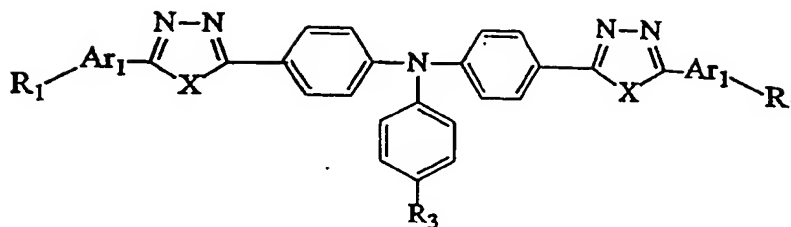
4. The OLED (10) of claim 2 or 3 wherein each R₁ is dimethylamino or diphenylamino.

5. The OLED (10) of one of claims 2 to 4 wherein each Ar₁ is 1,4-phenylene or 1,3-phenylene.

6. The OLED (10) of one of claims 2 to 5 wherein each R₂ is H.

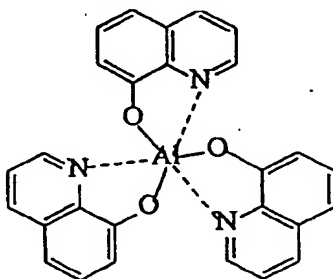
7. The OLED (10) of one of claims 2 to 6 wherein each R₃ is a C₁₋₂₀ alkyl, alkoxy, aryl, aryloxy, alkoxyaryl, or alkylcarbonyloxy group.

8. The OLED (10) of claim 1 wherein said oxadiazole compound has the formula

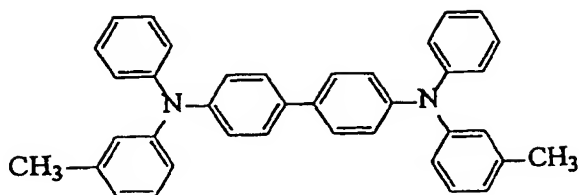


wherein X is independently in each occurrence -O-, -S-, or N(R₅), wherein R₅ is a C₁₋₂₀ hydrocarbyl (which correspond to the heterocyclic group being a 1,3,4 oxadiazole, 1,3,4-thiadiazole or 1,2,4,-triazole, respectively); R₁ is independently in each occurrence C₁₋₂₀ alkyl, alkoxy, aryl, aryloxy, or a tertiary amine group N(R₄)₂; R₃ is independently in each occurrence a C₁₋₂₀ alkyl, alkoxy group; and Ar₁ is independently in each occurrence a divalent C₅₋₁₀ aromatic group.

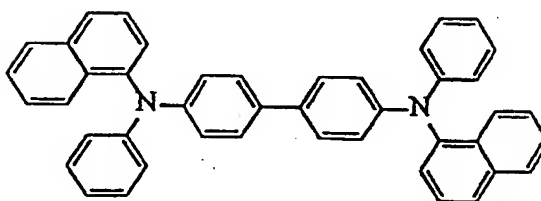
9. The OLED (10) of claim 8 wherein X is -O-, Ar₁ is 1,3-phenylene, R₁ is methoxy, and R₃ is methyl in each occurrence.
10. The OLED (10) of one of claims 1 to 9 wherein said cathode (16) comprises Mg or an alloy of Mg, Al or an alloy of Mg and Ag or an alloy of Li and Al.
11. The OLED (10) of one of claims 1 to 10 further comprising an electron transport layer (15), said electron transport layer (15) being sandwiched between said cathode (16) and said electroluminescent layer, said electron transport layer (15) comprising a compound having an energy band intermediate between that of said cathode (16) and that of said electroluminescent layer.
12. The OLED of claim 11 wherein said electron transport layer (15) comprises a metal chelated oxinoid compound and most preferably a tris(8-quinoline)Aluminium(Alq₃) whose formula is



13. The OLED (10) of claim 11 or 12 further comprising a hole transport layer (13), said hole transport layer (13) being sandwiched between said electroluminescent layer (14) and said anode (12), said hole transport layer (13) comprising a compound having an energy band intermediate between that of said anode (12) and that of said electroluminescent layer (14).
14. The OLED (10) of claim 13 wherein the hole transport compound is an aromatic tertiary amine.
15. The OLED (10) of one of claims 13 or 14 wherein said hole transport layer (13) comprises

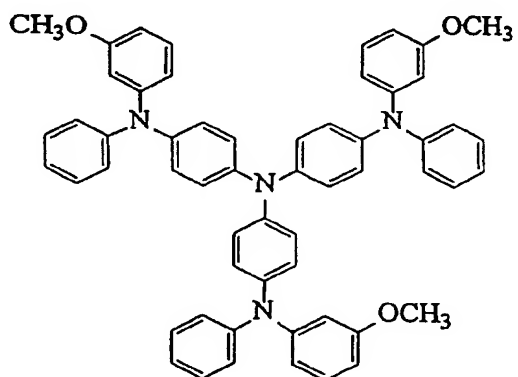


or



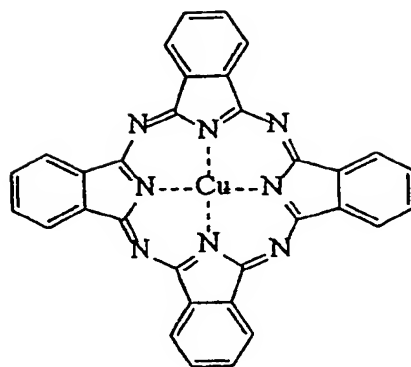
16. The OLED (10) of claim 13 wherein said anode (12) comprises first and second anode layers, said first anode layer (12) comprising indium tin oxide and said second anode layer (132) comprising a material that bonds to indium tin oxide and to said hole transport layer (131) and which has an energy band intermediate between that of indium tin oxide and said hole transport layer, said second anode layer being sandwiched between said hole transport layer and said first anode layer.

17. The OLED (10) of claim 16 wherein said second anode layer (132) comprises



or

18. The OLED (10) of claim 16 wherein said second anode layer (132) comprises



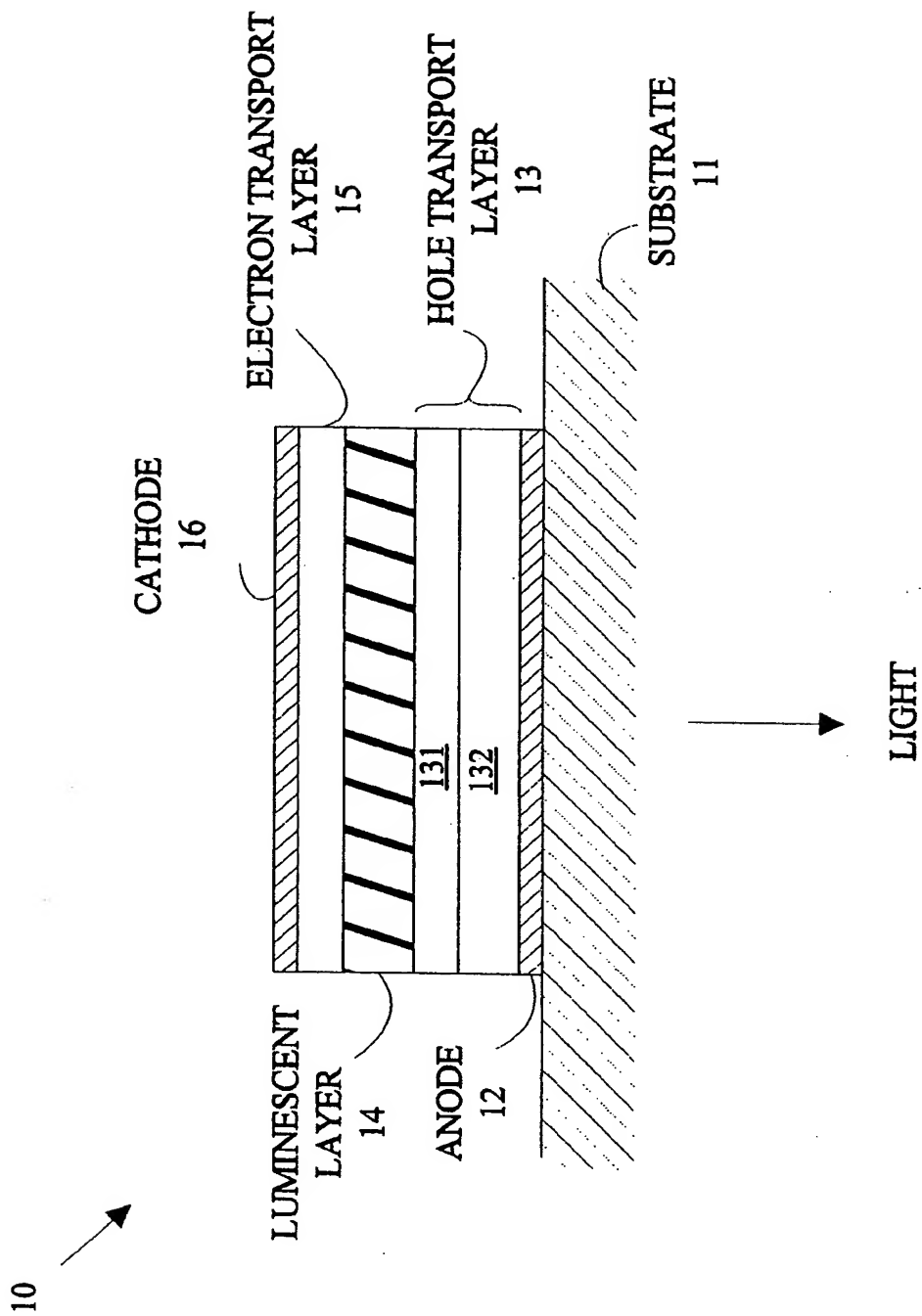


FIG. 1

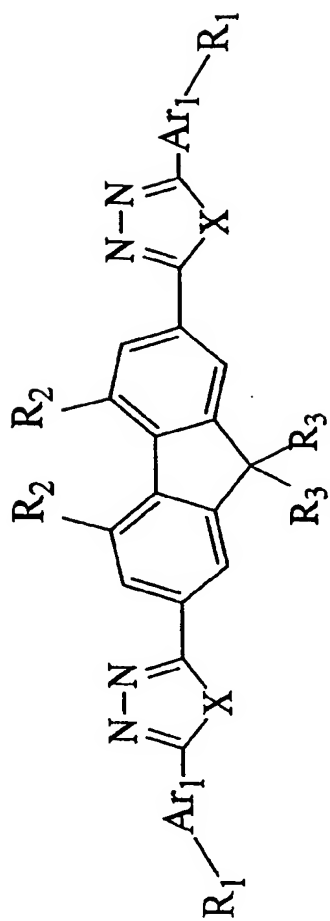


FIG. 2

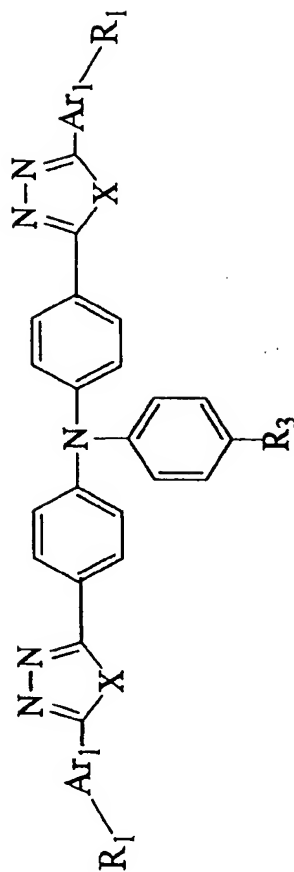


FIG. 3

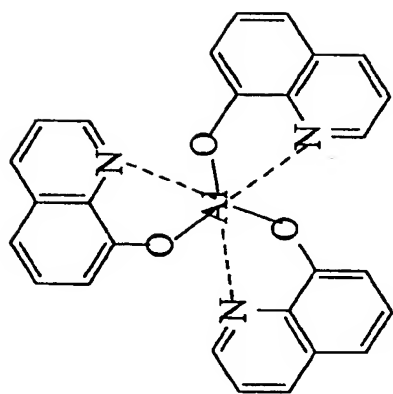


FIG. 4

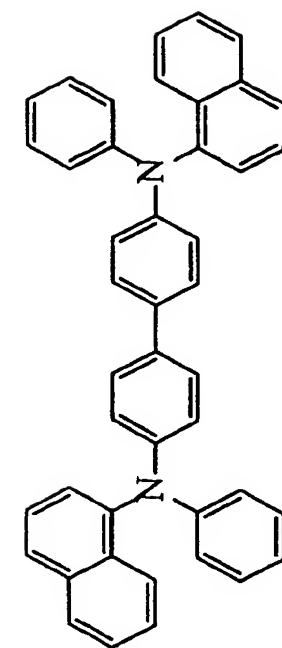


FIG. 6

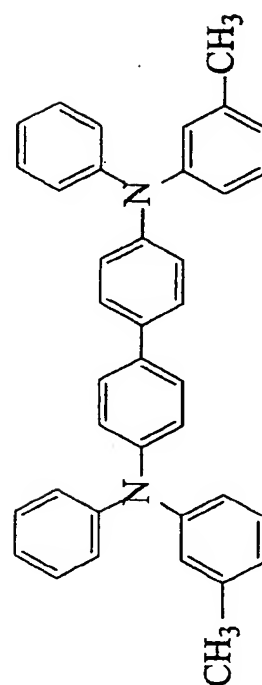


FIG. 5

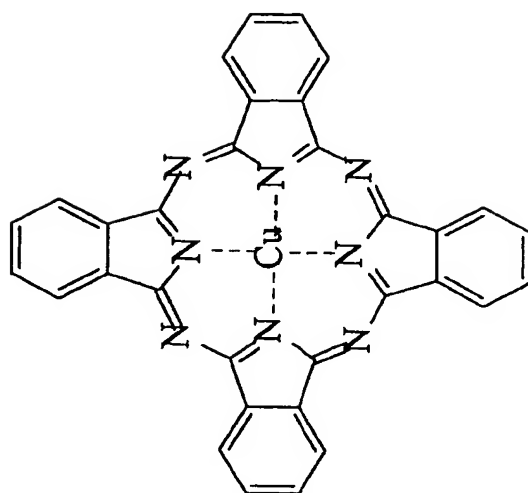


FIG. 8

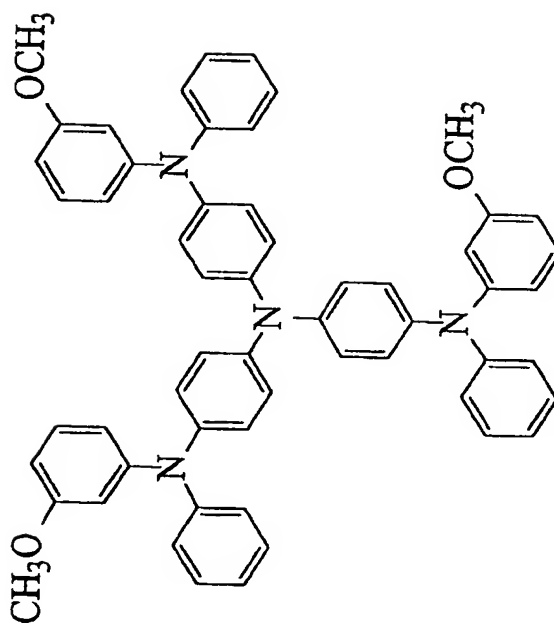


FIG. 7